Table 1. Analytical constituents for project. [Rank, H-high, M-medium, L-low]

Constituent		Analysis purpose	Rank(h,m,l)	Site purpose
	temperature, pH, conductance,			Standard data and allow for
	turbidity,	Standard purge monitoring parameters for sample stability		comparison to previous field
Field values	ORP, DO	criteria. Hydrogen ion activity and D.O. state.	н	parameter values.
	,	, ,		•
		Redox identification, iron available for oxide reactions. U+6		Not collected at site but easily
Dissolved ferrous iron	Fe2+	can be locked up in some cases with oxide formation.	Н	obtainable data.
Dissolved oxygen	DO	Additional measurement of D.O. as a check on probes.	Н	Check measurement on D.O. probe.
2.000 rea enjage.		Anion and cation balance. Understand carbonate and		enear measarement on 2101 proser
		bicarbonate reactions. U+6 mobile under high carbonate		Important constituent for charge
	Alkalinity as	concentrations in some cases. Alkaline leaching process		balance and to understand carbonate
Alkalinity	CaCO3	was employed at tailing sites.	Н	concentrations.
		General carbon availability gauge. Microbial-mediated		Limited data available at the site but
		reduction of uranium can be stimulated		provides useful information on
Total organic carbon	TOC	by the availability of an electron donor.	М	electron donor availability.
Majoranions	Dr. Cl. F. SO.4	Major anions for geochemical reactions and water type.	Н	Identify water type
Major anions	Br, Cl, F, SO4	iviajor amons for geochemical reactions and water type.	П	Identify water type.
Major cations	Ca, Mg, Na, K	Major cations for geochemical reactions and water type.	н	Identify water type.
,	224 trace	Includes important elements such as selenium and		· · · · · ·
Trace elements	elements	molybedenum; CO-CONTAMINANTS.	Н	Required constituents.
		High nitrate concentrations have been associated with the		
		site. Nitrate reactions can affect uranium concentrations.		
•		Presence of nitrate oxidized sulfur ores and previously		Understanding sources of nitrate is
Nitrogen	N, NO2 as N	immobile U+4 (PARADIS ET AL., 2016). Standard screening method for radioactivity; parameter	Н	important for assessing U mobility.
Gross alpha/beta		measured at site.	н	Required constituents.
p				,
	6 ll	Radium 226 is in the U-238 decay series. Radium 226 has a		
	Alpha- emitting	half life of 1600 years and decays into radon gas. Radium 228 is in the thorium-232 decay series. Radium has a half		
	•	life of 5.75 years. Radium 226 is a good indicator of waste		
	(226Ra and	water contamination from mines and mills (Kaufman et al.,		Identify natural and anthropogenic
Radium isotopes	228Ra)	1976).	н	contaminant.

		Radon 222 is a decay product of Radium 226. High radon- 222 has been associated with fault locations and uranium		
Radon-222	Radon-222	sources.	н	Radon-222 is a health hazard.
		There are three main isotopes of uranium (U-234, U-235, and U-238). U-238 is a weakly radioactive metal, and contributes to low-level background radiation in the environment. U-238 has a very long half-life of 4.47 billion years. Enriched U-235 is used as fuel in nuclear reactors and in nuclear weapons. Depleted uranium, which is poor in U-235 but rich in U-238, is used for commercial purposes. Uranium is common in specific types of igneous, metamorphic, and sedimentary rocks. Recent research indicates that increased concentrations of uranium in groundwater are caused by mobilization of uranium present in soil with irrigation waters containing bicarbonates. Also, nitrate can mobilize uranium through a series of bacterial and chemical reactions. Groundwater in the basin unaffected by mining has been found to have 234U/238U values ranging from 1.5 to 2.7. Ratios exceeding unity (secular equilibrium) result from alpha recoil of solid-phase 238U over geologic time causing the 234U daughter to be placed in favorable leaching sites. As ore bodies were dewatered, increased availability of oxygen caused uranium to dissolve rapidly without		Identify natural and anthropogenic
Uranium isotopes  Stable isotopes of water	U-238 2H/1H, 18O/16O	Assessment of recharge conditions, fractionation proceses such as evapotranspiration, and sources of water.	н	water affected by evaporation of tailings water from LTP can be identified.
Sulfur isotopes of sulfate	δ34S,δ32S	Sulfur reactions that are involved with U mobilization. Availability of pyrite and sulfur oxidation can impact U mobilization. These processes are visible with 32S and 34S ratios.	L	There is some evidence of gypsum availbility in geochemical reactions. Liberation of oxygen from dissolution of gypsum could enhance U mobilization.

Nitrogen isotopes	δ15N of NO3, δ14N of NO3, δ18O of NO3,δ16O of NO3 δ13C of DIC,	Theorectically, identify nitrate controls on U mobilization.  Dentrification processes can mobilize U and nitrogen isotopes can identify denitrification processes.	L	There are several sources of nitrate at the site including tailing operations and local septic systems.
Carbon isotopes  Carbon isotopes	δ13C of DOC, δ12C of DIC,δ12C of DOC	Useful for identifiying carbonate reactions and carbon dioxide influences on chemistry. Carbonate is an important factor in U mobilization. Carbon 14 has a large half life and is used to date very old waters. Groundwater in study area may contain a mix of old and new waters and this would help identify bimodal ages.	М	The alluvial system has carbonate waters that may promote U mobilization. Carbon isotopes may be useful for identifying mixing in Chinle.  Chinle waters may contain a mixture of old water.
Dissolved gases	N2, Ar, CO2, CH4, O2	Useful for looking at recharge conditions at time of water entering the subsurface.	L	Local and regional recharge may be distinguishable with measurements of dissolved gases. Measurements also provide additional check on oxygen and redox state.
GW dating: Tritium/Helium-3	tritium by	A primary age dating method for recennt (60 years) water. Peak TU is in late 1950's-early 1960's. Age is calculated from paren/daughter decay of <sup>3</sup> H and <sup>3</sup> He.	н	Selective wells be analyzed to assess recharge rates and time of travel. Understand formation of mixing zones in the Chinle and help tag U concentration to general ages of recharge.
CFCs	CFS	A primary age dating method for recent (40 years) water. Peak is in late 1980s.	н	Will be used as a check on tritium/helium ages at a subset of wells.
Helium-4	Helium, Neon, 3He/4He with tritium by 3He ingrowth	Screening tool to asses terrigenic Helium influence on tritium-helium dating.	М	Selectively analyzed constituent in Chinle waters.

1U.S. Environmental Protection Agency, 2009.

2Al, Sb, As, Ba, B, Be, Cd, Cr, Co, Cu, Fe, Li, Pb, Mn, Hg, Mo, Ni, Se, Si, Ag, Tl, U, V, and Zn.
3U.S. Environmental Protection Agency, 1993.
4{N2, Ar} R. F. Weiss, 1970, Deep-Sea Res., vol. 17, 721-735. R.F. {CO2} Weiss, 1974, Marine Chem. 2, 203-

215.[Bunsen Coef.]

{O2} B. B Beson and D. Krause,1980, Limnol. Oceanogr. 25(4) 662-671; 1984, Limnol. Oceanogr. 29(3), 620-632.

{CH4} D.A. Wiesenburg and N.L. Guinasso, 1979, J. Chem. Eng. Data Vol. 24, 356-360.



